

TEXAS DEPARTMENT OF HEALTH BUREAU OF RADIATION CONTROL

Application for Mammography Accreditation

Complete all items of the application in accordance with Title 25 Texas Administrative Code (TAC) §289.230(dd)(2). Please print or type. *Failure to furnish the requested information/documentation may result in a delay of the application review and/or disapproval of your application*. Retain a copy of the application for your files. Submit the appropriate fee(s) and the completed application to **either** address:

Postal service address:

Texas Department of Health Bureau of Radiation Control Texas Mammography Accreditation Program P.O. Box 149200 Austin, Texas 78756-3189

Overnight/express service or personal delivery address:

Texas Department of Health Bureau of Radiation Control-N127 Texas Mammography Accreditation Program 8407 Wall Street Austin, Texas 78754

If there are any questions, contact the Mammography Accreditation Program at (512) 834-6688.

Section 1: General Information			
Employer Identification Number (EIN):			
Legal Name of Facility:			
DBA(if applicable):			
Mailing Address:(Street/City/State/Zip)	Machine Use Location Address:(Street/City/State/Zip) (If multiple use locations, use additional sheets)		
Mammography Facility Phone Number:	FAX #:		
Lead Interpreting Physician:			
Telephone #:	E-mail address:		
	E-mail address:		
Section 2: Ty	pe of Action - Check all that apply		
" New Accreditation " Rene	ewal/Re-Instatement		
Who is your current accreditation body?	" N/A " ACR expiring on " State of Texas		
	Expiring:		
Do you wish to have your Certification expire the same month as your			

If you checked either N/A or State of Texas - skip to Section 4.

Section 3: Mammography Accreditation History

If you checked ACR in Section 2 you must complete this Section. Check one Yes "No 1. Are you requesting that the State of Texas be your accreditation body? "Yes "No 2. Provide the FDA Facility ID number assigned to the ACR accreditation: Expiration date: 3. When was the facility's current accreditation issued: 4. Was the accreditation ever denied? "Yes "No 5. Was the accreditation ever suspended or revoked? "Yes "No

- 6. If you checked **yes** to either question 4 or 5, you must provide a written explanation. (Use additional sheets) For question 4, you must provide the following:
 - a. Date(s) of failure(s);
 - b. Identity of the specific machine(s) that failed;
 - c. Details of the corrective action plan for the deficiencies, including required training for the physician, technologist and/or machine/processor repair;
 - d. Information on whether the machine(s) eventually passed the review after corrective actions were taken: and
 - e. Submit copies of the failure reports with this application.

For question 5, describe the circumstance that led to the suspension or revocation.

	Section 4: Facility Practice and Procedure Information			
1.	Type of Facility: check one	" Hospital" Hospital Outpatient Center" Clinic (specify type)	" Surgical Office er " Mobile (multiple locations)	
2.	Total number of machines:	Mammography units: Stereotactic Biopsy units:		
3.	Which of the following mammo " Screening	graphic procedures does your "Diagnostic	facility perform? <i>Check all that apply</i> "Self-Referral	
4.	. Average number of procedures performed per year at the facility:			
	Screening	Diagnostic	Aggregate (Total)	

Section 5: Personnel Qualifications

List all personnel involved with the mammography facility. For each individual, fill out the appropriate checklist and attach the required documentation to the form. Make copies of the forms as needed.

Interpreting Physician(s):		
Name:		
	,	
	•	
Radiologic Technologist(s) performing ma	mmography:	
Name:		
	•	
	•	
Medical Physicist(s): Name:		
	•	

Section 6: Mammography Equipment

Complete this section for each mammographic x-ray unit, except stereotactic units. The State of Texas does not accredit stereotactic biopsy units at this time. Check **all** appropriate boxes. Include a copy of a current medical physicist report for each machine. (Note - if there are any failures and/or deficiencies on the report, attach a list of corrective actions. Include copies of service/work invoices with the description of corrective actions.)

	" Located Onsite	Mammographic Unit <i>or</i>	" Mobile unit
1.	Control Panel Manufacturer:	Control Panel Model Name & Number:	Control Panel Serial Number:
2.	This machine is used for:	Screening "Diagnostic	" Magnification
3.	Target(s) & Filter(s) available:	Mo/Mo " Mo/Rh	" Rh/Rh
	Focal Spot Size(s):	W/Al " W/Mo ening: Magn	" W/Rh ification:
4.	Exposure Control:	Phototimed "Manual	
5.	Grid type used: Grid Ratio: enter a numeric ratio	Reciprocating "Stationary Grid frequency: er	nter a numberlines/cm or incl
6.	Type of imaging system: Screen/Film Combination:	Screen/Film " Digital en: Film:.	
7.	Analysis of Phantom Image: Technique used for phantom: Mode used:	ntom manufacturer and model number NP or AP "	mA & time
	Check all objects that are visi	ed on the phantom:	
	Fibers: " 1.56 millimeters " 1.12 millimeters " 0.89 millimeters " 0.75 millimeters	Specks: " 0.54 millimeters " 0.40 millimeters " 0.32 millimeters " 0.24 millimeters	Masses: " 2.00 millimeters " 1.00 millimeters " 0.75 millimeters " 0.50 millimeters
	0.54 millimeters0.40 millimeters	" 0.16 millimeters	" 0.25 millimeters

Must see entire fiber to count as a whole.

How many	specks in last
group?	

Must see rounded shape to count as a whole.

	Section 7:	Processing Equipment	
Manufacturer:	Dedicated Model N Mammography:	Number: Serial Nu	mber: Location:
	Yes " No		
	Yes " No		
If you have more	" Yes " No than one processor, please ind	icate which is the main	processor or the back-up processor(s).
Is batch processing If yes, subi	g utilized? "Yes "I mit film transport procedures {2		
	Section	on 8: Certification	
*Typed or printed		pplication is true and cu	Signature
Typed or printed	name of person who completed appli	cation Date	Signature
*This should be th facility.	ne signature of the Administrate	or, President, Chief Exec	rutive Officer, Owner or Partner of the
_	reting physician, I do hereby at attion with this application.	firm that I assume the res	ponsibilities listed in 25 TAC §289.230
Typed or printed	name of lead interpreting physician	Date	Signature

This MQSA Information Release Authorization must be signed by the Administrator, President, CEO, Owner, or Partner and the lead interpreting physician of the facility. Original signatures are required on this form. Stamps, electronic signatures or photocopied signatures are unacceptable.

MQSA Information Release Authorization

Effective October 1, 1994, all mammography facilities must be certified by the Food and Drug Administration (FDA), in accordance with the requirements of the Mammography Quality Standards Act (MQSA). The Texas Department of Health (TDH) has been designated by the FDA as one of the accreditation bodies in Texas for MQSA because its current accreditation program meets the standards contained in the MQSA law.

In order that facilities participating in the Texas Mammography Accreditation Program can obtain the required certification, the FDA will request the TDH to provide specific data about these facilities, units and personnel. The information would be drawn from the Application for Mammography Accreditation and other supporting documentation provided to TDH by the facility during the accreditation or re-accreditation process. Please complete the following authorization for release of information so that the necessary information may be forwarded to the FDA.

Authorization for Release of Information

"As the responsible parties of the mammography facility listed below, and in recognition of the need for this facility to be certified by the FDA, we hereby authorize the TDH to submit to the FDA any or all data about this facility that was submitted as part of its mammography accreditation or reaccreditation application, the survey report and findings made by the TDH, any other information about the facility that was submitted to the TDH or that the TDH currently has in its possession, and any other information gathered by the TDH in pre- or post-accreditation site visits by the TDH survey team."

Facility name and address:	
Signature of Administrator, President, CEO, Owner, or Partner	Signature of lead interpreting physician
Typed or printed name	Typed or printed name
Executed on	
Date	

Checklist for Interpreting Physicians

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You will need to make a copy of this form and use it for each in the requested document for each qualification listed below. At not send copies of DEA numbers, Controlled Substance licen	tach the required documentation to each form. Do
Initial qualifications:Texas Board of Medical Examiner's License - (cop	y of current license)
Date you started interpreting mammograms or qualifying date as established by MQSA inspector:	
quantying date as established by 11Q011 hispector.	(Month, Day, Year)

- 1. Based on the date stated above, select the proper category and submit the requested documentation. If you qualified prior to 04-28-1999, you must submit documentation of the following:
 - " ABR, AOBR or other certification body approved by FDA (copy of certificate), or Equivalent formal training, which includes two months of documented full-time training in interpretation of mammograms, including instruction in radiation physics, radiation effects and radiation protection. (Letter from the residency program director)
 - " *40 hrs formal training in mammography (self-attestation *or* letter from residency program director)
 - " *have read and interpreted 240 mammograms under the direct supervision of a qualified interpreting physician within a 6 month period. (self-attestation *or* letter from residency program director)

 If you qualified prior to 10-01-1994, direct supervision was not applicable.

Note: for those physicians who were in practice prior to October 1, 1994, a self-attestation will be accepted for those items with a * beside them. Self-attestations must state that these qualifications were met "prior to 10-01-1994". For physicians who completed their training after October 1, 1994, documentation must be submitted for all qualifications.

If you qualified after 04-28-1999, you must submit documentation of the following:

- " ABR, AOBR or other certification body approved by FDA (copy of certificate), *or* Equivalent formal training, which includes three months of documented full-time training in interpretation of mammograms, including instruction in radiation physics, radiation effects and radiation protection. (letter from the residency program director)
- " 60 hrs formal training in mammography (letter from residency program director)
- "If you passed your board at the first available opportunity, you will need to submit documentation that you have read and interpreted 240 mammograms under the direct supervision of a qualified interpreting physician within a 6 month period during the last 2 years of the residency program. (letter from residency program director) or

 If you did not take your board at the first available opportunity, you will need to submit documentation that you read and interpreted 240 mammograms under direct supervision of a qualified interpreting physician within the six month period prior to qualifying as an interpreting physician. (letter from supervising physician)

3. Continuing experience and education:

- " have read and interpreted mammograms for an average of 40 mammograms/month for 24 months (960 total) (Documentation of numbers from the facility are required, self-attestations are **not** allowed)
- " 15 continuing education units (CEUs) in **mammography** over a 3 year period (copies of mammography certificates, *do not send any other certificates*)

Checklist for Radiologic Technologist

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Name:	
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You will need to make a copy of this form and use it for each technologist at the facility who will be performing mammography. Please submit the requested document for each qualification listed below. Fill out the form completely and attach the requested documentation to each form.

1. Initial Qualifications:

" Medical Radiologic Technologist license - (copy of current license)

<u>Date you started performing mammography or</u>

qualifying date as established by MQSA inspector:

(Month, Day, Year)

2. Based on the date stated above, select the proper category and submit the requested documentation. If you qualified prior to 10-01-1994, you must submit documentation of the following:

- " 20 hours of formal mammography training; (letter from training program; CME certificate; documentation from in-house training program or self-attestation)
 - ARRT(M) (copy of card)

ard) Or

Date examination was taken: ______

If you qualified from 10-01-1994 to 10-01-1996, you must submit documentation of the following:

" 20 hours of formal mammography training; (letter from training program; CME certificate; or documentation from in-house training program)

Or

" ARRT(M) - (copy of card)

Date examination was taken:

If you qualified from 10-01-1996 to 08-10-1998, you must submit documentation of the following:

" 40 hours of formal mammography training; (letter from training program; CME certificate; documentation from in-house training program)

Note: During this time frame the FDA required 40 hours of formal training, but under certain circumstances accepted less. If you qualified with fewer hours, you must include a written statement that attests fewer hours were accepted by the MQSA inspector. Copies of the qualifying documentation must be submitted with the statement.

Or

" ARRT(M) - (copy of card)

Date examination was taken:

If you qualified from 08-10-1998 to 04-28-1999, you must submit documentation of the followin:

" 40 hours of formal mammography training; (letter from training program; CME certificate; documentation from in-house training program)

If you qualified after 04-28-1999 you must submit documentation of the following:

- " 40 hours formal training in mammography; (letter from training program; CME certificate; documentation from in-house training program)
- " performed 25 mammograms under direct supervision (letter from training program; documentation from in-house training program)

3. Continuing experience and education:

' 15 CEUs in mammography over a 36 month period (copies of certificates)

As of April 28, 1999 you must document the performance of 200 mammograms over 24 months. Documentation of numbers from facilities will be required; **self-attestations are not accepted**. (Inspectors will start reviewing records 04-28-2001.

Checklist for Medical Physicist

Name:			
	·		

You will need to make a copy of this form and use it for each medical physicist at the facility who will be performing the annual mammography system survey. Please submit the requested document for each qualification listed below. Fill out the form completely and **attach** the requested documentation to each form.

1. " Texas Medical Physics Practice Act license - (copy of current license)

Qualifying date as established by MQSA inspector:

(Month, Day, Year)

2. Select the proper category and submit the requested documentation.

If you qualified under the Initial Qualifications, you must submit documentation of the following:

- " Master's degree or higher in a physical science (copy of degree)
- " 20 semester hours in physics (copy of college transcript or letter from college stating hours)

 Note: If the degree is in physics, this documentation will not need to be submitted.
- " 20 contact hours of specialized training in surveying mammography equipment
- " experience conducting surveys one mammography facility & 10 mammography units

<u>If you qualified under the Alternative Initial Qualifications, prior to April 28, 1999, you must submit the following documentation:</u>

- " Bachelor's degree or higher in a physical science (copy of degree)
 - Note: training and experience must be met after fulfilling degree requirements.
- " 10 semester hours in physics (copy of college transcript or letter from college stating hours)

 Note: If the degree is in physics, this documentation will not need to be submitted.
- " 40 contact hours of specialized training in surveying mammography equipment
- " experience conducting surveys one mammography facility & 20 mammography units

3. Continuing experience and education:

" 15 CEUs in over a 36 month period (copies of certificates)

As of April 28, 1999 you must document the performance of surveys of a total of two mammography facilities and six mammography units over 24 months. Documentation of surveys will be required; **self-attestations are not accepted**. (Inspectors will start reviewing records 04-28-2001.)



FEES FOR ACCREDITATION AND/OR CERTIFICATION OF MAMMOGRAPHY FACILITY

Each new application for accreditation and/or certification of a mammography facility shall be accompanied by the appropriate fees. In addition, each renewal accreditation application shall be accompanied by the appropriate fees. No application will be accepted for filing or processed prior to payment of the full amount due. [25 TAC §289.230 (ee) and §289.204 (h)]

The fee(s) for accreditation of your mammography facility will be one or more of the following:					
	"	Accreditation for the first mammography unit	\$ 720.00		
	"	Accreditation fee for each additional mammography unit (Number of additional mammography unit(s) x \$345.00)			
		ACCREDITATION TOTAL D	DUE	<u>\$</u>	
,,	Th	e fee for certification of your mammography facility will be:			
	"	\$422.00 per mammography unit (number of mammography unit(s) x \$422.00)	\$		
		CERTIFICATION TOTAL D	UE	<u>\$</u>	
		TOTAL DUE WITH APPLIC	ATION(S)	\$	
ma	nne	complete this form and submit it with your application so r. If you have any questions regarding the payment of these fees, y ation Control at (512) 834-6688.		-	
Na	me	of facility:			

http://www.tdh.state.tx.us/ech/rad/pages/brc.htm An Equal Employment Opportunity Employer